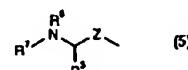
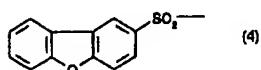
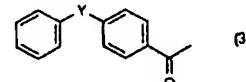
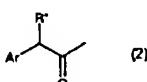
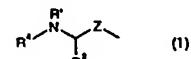
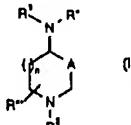




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(57) Abstract			
<p>This invention relates to compounds of formula (I), wherein A is C(O) or CH(OH); R¹ is (1), (2), (3) or (4); R² is H, C₁-6alkyl, C₃-6cycloalkyl-C₀-6alkyl, Ar-C₀-6alkyl, Het-C₀-6alkyl, R⁵C(O)-, R⁵C(S)-, R⁵SO₂-, R⁵OC(O)-, R⁵R'NC(O)-, R⁵R'NC(S)-, adamantly-C(O)-, or (5); R" is H, C₁-6alkyl, Ar-C₀-6alkyl, or Het-C₀-6alkyl; R'" is H, C₁-6alkyl, C₃-6cycloalkyl-C₀-6alkyl, Ar-C₀-6alkyl, or Het-C₀-6alkyl; each R³ independently is H, C₂-6alkenyl, C₂-6alkynyl, Het, Ar or C₁-6alkyl optionally substituted by OR¹, SR¹, NR¹2, R'NC(O)OR⁵, CO₂R¹, CO₂NR¹2, N(C=NH)NH₂, Het or Ar; R⁴ is H, C₁-6alkyl, C₃-6cycloalkyl-C₀-6alkyl, Ar-C₀-6alkyl, Het-C₀-6alkyl, R⁵C(O)-, R⁵C(S)-, R⁵SO₂-, R⁵OC(O)-, R⁵R'NC(O)-, R⁵R'NC(S)-, R'HNCH(R')C(O)-, or R⁵OC(O)NR'CH(R')C(O)-; each R⁵ independently is C₃-6cycloalkyl-C₀-6alkyl, Ar-C₀-6alkyl, Het-C₀-6alkyl, Ar-C₀-6alkoxy, Het-C₀-6alkoxy, or C₁-6alkyl optionally substituted by OR¹, SR¹, NR¹2, R'NC(O)OR⁵, CO₂R¹, CO₂NR¹2, N(C=NH)NH₂, Het or Ar; R⁶ is H, C₁-6alkyl, Ar-C₀-6alkyl, or Het-C₀-6alkyl and R⁷ is H, C₁-6alkyl, C₃-6cycloalkyl-C₀-6alkyl, Ar-C₀-6alkyl, Het-C₀-6alkyl, R⁵C(O)-, R⁵C(S)-, R⁵SO₂-, R⁵OC(O)-, R⁵R'NC(O)-, R⁵R'NC(S)-, R'HNCH(R')C(O)-, or R⁵OC(O)NR'CH(R')C(O)-; or R⁶ and R⁷ are connected to form a pyrrolidine, a piperidine, or a morpholine ring; each R' independently is H, C₁-6alkyl, Ar-C₀-6alkyl, or Het-C₀-6alkyl; R* is H, C₁-6alkyl, C₃-6cycloalkyl-C₀-6alkyl, Ar-C₀-6alkyl, or Het-C₀-6alkyl; Y is a single bond or O; each Z independently is CO or CH₂; and n is 0, 1, or 2; or a pharmaceutically acceptable salt thereof, which are inhibitors of cysteine proteases, particularly cathepsin K, and are useful in the treatment of diseases in which inhibition of bone loss is a factor.</p>			



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